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(71) Applicant: Halliburton Energy Services, Inc. Duncan, Oklahoma 73536 (US)

(72) Inventors:

Vijn, Jan Pieter
 2352 VV Leiderdorp (NL)

Dao, Bach
 2151 KK Nieuw Vennep (NL)

(74) Representative: Waln, Christopher Paul et al
 A.A. Thornton & Co.
 235 High Hoiborn
 London WC1V 7LE (GB)

(54) Water-microsphere suspensions for use in well cements

(57) Storable water-microsphere suspensions for use in well cements are basically comprised of water, microspheres and an effective amount of a suspending agent selected from microfine or colloidal materials and

gel forming polymers. The suspensions are mixed with a well cement composition, and the composition is then introduced into a well and allowed to set.

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Description

[0001] The present invention relates to storable water-microsphere suspensions and their use in well cements.

[0002] Hydraulic cement compositions are commonly utilized in subterranean well completion and remedial operations. For example, hydraulic cement compositions are used in primary cementing operations whereby strings of pipes such as casings and liners are cemented in well bores. In performing primary cementing, a hydraulic cement composition is pumped into the annular space between the walls of the well bore and the exterior surfaces of a pipe string disposed therein. The cement composition is permitted to set in the annular space thereby forming an annular sheath of hardened substantially impermeable cement therein. The cement sheath physically supports and positions the pipe string in the well bore and bonds the exterior surfaces of the pipe to the walls of the well bore whereby the undesirable migration of fluids between zones or formations penetrated by the well bore is prevented.

[0003] In some locations, the subterranean zones or formations into or through which wells are drilled have high permeabilities and low compressive and tensile strengths. As a result, the resistances of the zones or formations to shear are low and they have very low fracture gradients. When a well fluid such as a hydraulic cement composition is introduced into a well bore penetrating such a subterranean zone or formation, the hydrostatic pressure exerted on the walls of the well bore can exceed the fracture gradient of the zone or formation and cause fractures to be formed in the zone or formation into which the cement composition is lost. While lightweight cement compositions have been developed and used which contain microspheres for reducing the density of the cement compositions, the microspheres are usually dry blended with the cement utilized followed by mixing the dry blend with water. This procedure, however, is troublesome because it is difficult to obtain a constant cement-microsphere blend due to the differences in density between the cement and microspheres. While the microspheres can be added to the mix water, because the microspheres are lighter than the water, they float in the water.

[0004] Thus, there are needs for storable water-microsphere suspensions for use in preparing lightweight well cement compositions which can be stored at the well site or on an offshore platform for use when needed.

[0005] In one aspect, the invention provides a storable water-microsphere suspension for use in well cement compositions comprising water microspheres and a suspending agent selected from microfine or colloidal materials and

[0006] In another aspect, the invention provides a method of cementing a well bore, which method comprises the

(a) combining a water-microsphere suspension of the invention with a cement composition comprised of cement

(b) introducing said cement composition including said water-microsphere suspension into said well bore; and

(c) allowing said cement composition to set.

[0007] The storable water-microsphere suspensions of the invention can be used for forming lightweight well cement compositions. While being stored, the suspensions normally do not separate for at least three weeks, and normally do not require stirring or agitation before use.

[0008] As mentioned above, microspheres have heretofore been blended with cement compositions to reduce the density of the cement compositions. The microspheres have generally been dry blended with the cement. However, it is often difficult to obtain a uniform microsphere-cement mixture due to the particle size and density difference of the spheres as compared to the cement particles. An alternate method of combining the microspheres with a cement composition is to add the microspheres to the mix water before combining the cement therewith. However, because the microspheres are lighter than water they float in the water making it difficult to achieve a mixture.

[0009] The present invention provides stabilized water-microsphere suspensions which are highly resistant to separation, can be stored at the job site in containers for three weeks or longer and can be added to the cement composition mixing water or to the cement composition after it has been mixed. The density of the final cement composition can be easily adjusted by adding more or less of the suspension to the cement composition. The storage of the homogenous water-microsphere suspension at the well site allows the suspension to be added to a cement composition in the required amount and any of the water-microsphere suspension not utilized can be re-stored and utilized later on another

[0010] The water utilized to form a storable water-microsphere suspension of this invention is preferably fresh water or salt water. The term "salt water" is used herein to mean unsaturated salt solutions and saturated salt solutions including brine and seawater. Generally, water from any source can be utilized so long as the water does not react with any of the components in the cement composition to which the water-microsphere suspension is added.

[0011] The microspheres utilized in the water-microsphere suspensions of this invention can be fly ash microspheres, glass microspheres or recycled glass microspheres. Of these, fly ash microspheres are preferred.

[0012] Particularly suitable fly ash microspheres are commercially available from Halliburton Energy Services, Inc.

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of Duncan, Oklahoma, under the tradename "SPHERELITETM". Synthetic hollow glass microspheres are commercially available from the Minnesota, Mining and Manufacturing Company under the tradename "SCOTCHLITETM". The microspheres utilized are included in a suspension of this invention in an amount sufficient to produce a suspension density in the range of from about 4.6 to about 9.5 pounds per gallon, i.e., in an amount in the range of from about 25% to about 150% by weight of the water in the suspension.

[0013] The suspending agent in the water-microsphere suspension functions to prevent separation of the microspheres from the water and to maintain a homogenous suspension while being stored. In accordance with the present invention, the suspending agent is selected from microfine or colloidal materials or gel forming polymers. Examples of microfine or colloidal materials which can be used include, but are not limited to, carbon black, lignite, brown coal, humic acid, fumed silica, precipitated silica, polyvinyl alcohol latex, styrene-butadiene latex and surfactant micelle. Of these, carbon black is presently preferred. The polymers which can be used include, but are not limited to, carragenan, scleroglycan, xanthan, guar, hydroxypropylguar, hydroxyethylcellulose, carboxymethylhydroxyethylcellulose and copolymers or terpolymers of acrylamidemethyl propylene sulfonate, N,N-dimethylacrylamide and acrylic acid. Of these, carragenan is preferred.

[0014] One or more of the above described suspending agents are generally included in the water-microsphere suspension in an amount in the range of from about 0.1% to about 75% by weight of the water in the suspension. When the suspending agent is a microfine or colloidal material as described above, the microfine or colloidal material is preferably included in the water-microsphere suspension in an amount in the range of from about 25% to about 75% by weight of water therein, more preferably in an amount of about 49%. When the suspending agent is a polymer described above, the polymer is preferably included in the water-microsphere suspension in an amount in the range of from about 0.1% to about 1.5% by weight of water therein, more preferably in an amount of about 0.5%.

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[0015] The methods of cementing a well bore of the present invention are basically comprised of the following steps. A water-microsphere suspension for reducing the density of a well cement composition comprised of water, microspheres and an effective amount of a suspending agent selected from the group of microfine or colloidal materials and gel forming polymers is stored. Thereafter, the water-microsphere suspension is combined with a cement composition comprised of a hydraulic cement and water. The cement composition including the water-microsphere suspension is introduced into a well bore and the cement composition is allowed to set therein.

[0016] The hydraulic cements eutilized in the cement composition can be those comprised of calcium, aluminum, silicon, oxygen and/or sulfur which set and harden by reaction with water. Such hydraulic cements include Portland cements, pozzolana cements, gypsum cements, high aluminum content cements, silica cements, high alkalinity cements and slag cements. The cements can be of conventional particle sizes or they can be of ultra-fine particle sizes. Portland cements are generally preferred for use in accordance with this invention. Portland cements of the types defined and described in API Specification For Materials And Testing For Well Cements, API Specification 10, 5th Edition, dated July 1, 1990 of the American Petroleum Institute are particularly suitable. Preferred such API Portland cements include classes A, B, C, G and H, with API classes G and H being more preferred and class G being the most preferred. Portland cements of the types defined and described in European Norms 197-1:2000 Cement - Part 1: Composition, specifications and conformity criteria for common cements CEN publication date 2000-06-21 are also preferred. The cements are classified as CEM I portland cement-possolan mixtures, and CEM V composite cement blends.

[0017] The water utilized in the cement composition can be fresh water or salt water. As mentioned above, the term "salt water" is used herein to mean unsaturated salt solutions and saturated salt solutions including brines and seawater. The water is generally present in the cement composition in an amount sufficient to form a pumpable slurry, generally an amount in the range of from about 38% to about 56% by weight of cement in the composition. The water-microsphere suspension utilized in accordance with the methods of this invention is generally included in the cement composition in an amount to produce a cement composition density in the range of from about 9.8 to about 15.0 pounds per gallon, i.e., in an amount in the range of from about 20% to about 300% by weight of water in the cement composition.

[0018] A preferred storable water-microsphere suspension of this invention for use in well cement compositions comprises: water; microspheres selected from the group consisting of fly ash microspheres, glass microspheres and recycled glass microspheres present in the suspension in an amount in the range of from about 25% to about 150% by weight of water therein; and a microfine or colloidal suspending agent selected from the group consisting of carbon black, lignite, brown coal, humic acid, fumed silica, precipitated silica, polyvinyl alcohol latex, styrene-butadiene latex and surfactant micelle present in the suspension in an amount in the range of from about 25% to about 75% by weight of water therein. The most preferred microfine or colloidal suspending agent for use in the above described suspension is carbon black.

[0019] Another preferred storable water-microsphere suspension of this invention for use in well cement compositions comprises: water; microspheres selected from the group consisting of fly ash microspheres, glass microspheres and recycled glass microspheres present in the suspension in an amount in the range of from about 25% to about

150% by weight of water therein; and a polymer suspending agent selected from the group consisting of carragenan, scleroglycan, xanthan, guar, hydroxypropylguar, hydroxyethylcellulose, carboxymethylhydroxyethylcellulose and copolymers or terpolymers of acrylamidemethyl propylene sulfonate, N,N-dimethylacrylamide and acrylic acid present in the suspension in an amount in the range of from about 0.1% to about 1.5% by weight of water therein. The most preferred polymer suspending agent for use in the above described suspension is carragenan.

[0020] A preferred method of cementing a well bore of this invention comprises the steps of: (a) storing a water-microsphere suspension for reducing the density of a well cement composition comprised of water, microspheres selected from the group consisting of fly ash microspheres, glass microspheres and recycled glass microspheres present in the suspension in an amount in the range of from about 25% to about 150% by weight of water therein and a microfine or colloidal suspending agent selected from the group consisting of carbon black, lignite, brown coal, humic acid, fumed silica, precipitated silica, polyvinyl alcohol latex, styrene-butadiene latex and surfactant micelle present in the suspension in an amount in the range of from about 25% to about 75% by weight of water therein; (b) combining the water-microsphere suspension with a cement composition comprised of a hydraulic cement and water; (c) introducing the cement composition including the water-microsphere suspension into the well bore; and (d) allowing the cement composition to set.

[0021] Another preferred method of cementing a well bore of the present invention comprises the steps of: (a) storing a water-microsphere suspension for reducing the density of a well cement composition comprised of water; microspheres selected from the group consisting of fly ash microspheres, glass microspheres and recycled glass microspheres present in the suspension in an amount in the range of from about 25% to about 150% by weight of water therein and a polymer suspending agent selected from the group consisting of carragenan, scleroglycan, xanthan, of acrylamidemethyl propylene sulfonate, N,N-dimethylacrylamide and acrylic acid present in the suspension in an amount in the range of from about 0.1% to about 1.5% by weight of water therein; (b) combining the water-microsphere suspension with a cement composition comprised of a hydraulic cement and water; (c) introducing the cement composition including the water-microsphere suspension into the well bore; and (d) allowing the cement composition to set. [0022] In order to further illustrate the storable water-microsphere suspensions and methods of cementing in a well bore in accordance with the present invention, the following example is given.

EXAMPLE

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[0023] Glass microspheres were suspended in fresh water containing 49% microfine carbon black by weight of the water to produce a suspension containing 89% glass microspheres by weight of the water. The suspension, identified as Sample No. 1, was stored and monitored over a period of 3 weeks. That is, the sample was examined for separation of the microspheres whereby they float on top of the water. If separation occurs, the microspheres form a cake on top of the suspension which prevents pouring or pumping of the suspension. Also, the development of free water and settling and whether the suspension was re-stirrable to its original consistency were monitored after 3 days and after 3 weeks. The results of these test are given in Table I below.

[0024] Fly ash microspheres were also suspended in fresh water containing 0.5% carragenan polymer by weight of the water to produce a suspension containing 50% fly ash microspheres by weight of the water. The suspension, identified as Sample No. 2, was also stored and monitored over a period of 3 weeks as described above for Sample No. 1. The results of these tests are also given in Table I below.

[0025] Portions of Sample No. 1 and Sample No. 2 suspensions were added to portions of a cement composition comprised of CEM I 52.5 Portland hydraulic cement according to EN 197, fresh water in an amount sufficient to form a slurry, and a dispersing agent comprised of the condensation product of acetone, formaldehyde and sodium sulfite present in an amount of about 1.0% by weight of the hydraulic cement. The portions of the Sample No. 1 and Sample No. 2 suspensions were present in the cement composition portions in amounts of 117% and 120%, respectively, by weight of water in the cement composition portions.

[0026] Samples of the above described cement compositions were tested for rheologies and densities at room temperature and pressure, at 4,000 psi and at 125°F. The cement compositions were also tested for free water, settling, top/bottom densities and compressive strengths at 125°F using a Universal Cement Analyzer. The results of these tests are given in Table II below.

TABLE I

	Stability O	Microsphere Suspensions				
5	Suspension Sample Components, Properties And Stability	z Suspension Sample No.				
		1	2			
	Water					
10	Carbon Black, % by wt. of water	49	-			
	Carragenan, % by wt. of water	-	0.5			
	Glass Microspheres, % by wt. of water	89	-			
	Fly Ash Microspheres, % by wt. of water	•	50			
15	Density At Atmospheric Pressure, lb/gal	7.15	7.33			
	Density At 4,000 psi, lb/gal	7.15	8.25			
	Rheology 300-200-100-6-3	131-101-66-19-15	105-90-71-35-3			
	Stability:					
20	After 3 days	no separation and free flowing	no separation and free flowing			
20	After 3 weeks	no separation and free flowing	no separation and free flowing			

TABLE II

25	Properties Of Cement Comp	ositions Containing Microspher	re Suspensions
	Test Cement Sample Components And Properties	Cement Compositions Containing	g Test Microsphere Suspensions
	Portland cement CEM 52.5		
30	Water, % by wt. of cement	44	44
	Dispersant ¹ , % by wt. of cement	1.0	None
	Sample No. 1 Suspension, % by wt. of water	117	None
	Sample No. 2 Suspension, % by wt. of water	None	120
35	Density At Atmospheric Pressure, lb/gal	12.0	11.9
	Density At 4,000 psi, lb/gal	12.1	12.5
40	Total Amount Of Water In Cement Composition, % by wt. of cement	63.3	61.6
	Microsphere Flotation After Adding Test Suspension To Mix Water	None	None
45	Rheology At Atmospheric Temperature And Pressure, 300-200-100-60-30-6-3	87-69-49-40-32-24-23	30-23-14-11-8.5-7-6.5
	Density, lb/gal	12.0	11.9
50	Rheology At 4,000 psi, 300-200-100-60-30-6-3	87-69-50-42-34-25-24	53-42-30-25-21-17-15
	Density, lb/gal	12.1	12.5
	Rheology At 125°F, 300-200-100-60-30-6-3	79-62-44-36-28-20-19	62-54-45-41-36-20-15
55	Density, lb/gal	12.0	12.5

¹Condensation product of acetone, formaldehyde and sodium sulfite.

TABLE II (continued)

	Properties Of Cement Co	ompositions Containing Microsphere	9 Suspensions			
5	Test Cement Sample Components And Properties	Cement Compositions Containing Test Microsphere Suspensions				
	Portland cement CEM 52.5					
0	Freewater, % by vol. Settling, % by vol. Density, 1b/gal	· -	0.4 0 Gelled			
5	Compressive Strength ² , at 125°F 50 psi at hrs:min 500 psi at hrs:min psi at 24 hrs psi at 72 hrs Crush, psi Cube Crush - 7 days, psi	6:24 6:53 3750 4634 4679 4688	1:56 2:48 2275 2600 4297			

²Universal Cement Analyzer.

[0027] From Tables I and II, it can be seen that the microsphere suspensions of the present invention can be stored for at least 3 weeks while remaining stable and can be combined with hydraulic cement compositions without adverse

Claims

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- 1. A storable water-microsphere suspension for use in well cement compositions comprising water microspheres and a suspending agent selected from microfine or colloidal materials and gel forming polymers.
 - 2. A suspension according to claim 1, wherein the water is fresh water or salt water.
- 3. A suspension according to claim 1 or 2, wherein the microspheres are fly ash microspheres, glass microspheres 35 or recycled glass microspheres.
 - 4. A suspension according to claim 1, 2 or 3, wherein said microspheres are present in said suspension in an amount of from 25% to 150% by weight of water in said suspension.
 - 5. A suspension according to claim 1, 2, 3 or 4, wherein said suspending agent is a microfine or colloidal material selected from carbon black, lignite, brown coal, humic acid, fumed silica, precipitated silica, polyvinyl alcohol latex, styrene-butadiene latex and surfactant micelle; or the suspending agent is a polymer selected from carragenan, scleroglycan, xanthan, guar, hydroxypropylguar, hydroxyethylcellulose, carboxymethylhydroxyethylcellulose and a copolymer or terpolymer of acrylamidemethyl propylene sulfonate, N,N-dimethylacrylamide and acrylic acid.
 - 6. A suspension according to any of claims 1 to 5, wherein said suspending agent is present in said suspension in an amount of from 0.1% to 75% by weight of said water in said suspension.
- 7. A suspension according to claim 1 or 2, wherein the microspheres are fly ash microspheres, glass microspheres 50 or recycled glass microspheres and are present in said suspension in an amount of from 25% to 150% by weight of water therein; and the microfine or colloidal suspending agent is selected from carbon black, lignite, brown coal, humic acid, fumed silica, precipitated silica, polyvinyl alcohol latex, styrene-butadiene latex and surfactant micelle, present in said suspension in an amount of from 25% to 75% by weight of water therein. 55
 - 8. A suspension according to claim 1 or 2, wherein the microspheres are selected from fly ash microspheres, glass microspheres and recycled glass microspheres and are present in said suspension in an amount of from 25% to 150% by weight of water therein; and the suspending agent is a gel forming polymer selected from carragenan,

scleroglycan, xanthan, guar, hydroxypropylguar, hydroxyethylcellulose, carboxymethylhydroxyethylcellulose and a copolymer or terpolymer of acrylamidemethyl propylene sulfonate, N,N-dimethylacrylamide and acrylic acid, present in said suspension in an amount of from 0.1% to 1.5% by weight of water therein.

- 5 9. A method of cementing a well bore, which method comprises the steps of
 - (a) combining a water-microsphere suspension, as claimed in any of claims 1 to 8, with a cement composition comprised of cement and water;
 - (b) introducing said cement composition including said water-microsphere suspension into said well bore; and
 - (c) allowing said cement composition to set.

10. A method according to claim 9, wherein the water-microsphere suspension is stored prior to use.



EUROPEAN SEARCH REPORT

Application Number

EP 02 25 7612

	DOCUMENTS CONSID	ERED TO BE RELEVA	NT	
Category	Citation of document with in of relevant passa	dication where specials	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.CI.7)
X	EP 0 748 782 A (SOF 18 December 1996 (1 * the whole documen	996-12-181	1-8	E21B33/13
(US 6 153 562 A (BAR AL) 28 November 200 * example 1 *	ET JEAN-FRANCOIS E 0 (2000-11-28)	T 1-8	
	WO 93 09332 A (ATLA 13 May 1993 (1993-09 * examples *	NTIC RICHFIELD CO) 5-13)	1-8	
1	WO 98 38855 A (ENGE AGRICULTURE (US)) 11 September 1998 (1 * claims 1,9 *		1-3,6	
[]	EP 0 492 087 A (OPTI KUNSTSTOFFS) 1 July * examples *	PLAST GES FUER 1992 (1992-07-01)	1,2,6,7	
1 /	DD 296 474 A (VE ING AUTOM) 5 December 19 * claims 3,4 *	ENIEURBETRIEB FUER 91 (1991-12-05)	1-3,8	TECHNICAL FIELDS SEARCHED (Int.CI.7) E218 C04B
! 1	/S 4 623 390 A (DELM .8 November 1986 (196 examples *	 ONICO JOHN A) 86-11-18)	1,2	
Tr	ne present search report has beer	denum in to the state of		
	ace of search			
	INICH	Date of completion of the search 7 March 2003	Polli	Examiner O M
: particular : particular documen : technolog	SORY OF CITED DOCUMENTS rly relevant if taken alone rly relevant if combined with another t of the same category gical background en disclosure	T: theory or princ E: earlier patent of after the filing D: dooument cite L: dooument cite	iple underlying the inver	ition On, or

EPO FORM 1503 03.82 (P04C01)

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO.

EP 02 25 7612

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

07-03-2003

Patent docume cited in search re		Publication date		Patent fam member(s		Publication date
EP 0748782	A	18-12-1996	FR	2735465	A1	20-12-1990
			CA	2178707		14-12-1990
			ΕP	0748782		18-12-1990
			ĞB	2302091		08-01-1997
			ÑÕ	962474		16-12-1990
			ÜS	6312515		06-11-200
US 6153562	Α	28-11-2000	FR	2753963	A1	03-04-1998
			ΑU	3922397	Α	02-04-1998
			CA	2216750	A1	30-03-1998
			EΡ	0832861	A1	01-04-1998
			NO	974528	Α	31-03-1998
			ΑU	731607	B2	05-04-200
			DE	69705475	D1	09-08-200
			DE	69705475	T2	25-04-200
			DK	832861	. –	15-10-200
WO 9309332	Α	13-05-1993	US	5211238	Α	18-05-1993
			US	5188176		23-02-1993
			WO.	9309332	A1	13-05-1993
WO 9838855	Α	11-09-1998	US	5908708		01-06-1999
			AU	731684		05-04-200
			ΑU	6343598	• •	22-09-1998
			AU	731843		05-04-200
			AU	6443798	• -	22-09-1998
			AU	747461		16-05-2002
			AU	6443898		22-09-1998
			AU	6675398		22-09-1998
			BR	9808153	A	28-03-2000
			BR		A	28-03-2000
			BR	9812287	A	18-07-2000
			CN	1249663	Ţ	05-04-2000
			CN	1249664	Ţ	05-04-2000
			CN	1265003	Ţ	30-08-2000
			EP	1011309		28-06-2000
			EP	0969718		12-01-2000
			EP	0971593	A1	19-01-2006
			EP	0969725		12-01-2000
			IL	131474	Α	28-01-2001
			JP	2001513818	T	04-09-2001
			JP	2001524945	T	04-12-2001
			JP	2001524946	T	04-12-2001
			JP	2002515760	T	28-05-2002
			NZ	337203		27-03-2000
			ΝZ	337204	A	01-03-2002

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO.

EP 02 25 7612

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

07-03-2003

Patent de cited in sea	ocument rch report	Publication date		Patent family member(s)		Publication date
WO 983885	5 A		NZ TR TR TR WO WO WO US US US US US US ZA ZA ZA	337205 A 9902119 T 9902120 T 9902136 T 445133 B 9838866 A 9838855 A 9838867 A 9838848 A 2003020975 A 6060521 A 6069112 A 6235683 B1 6464995 B1 6156327 A 6010867 A 9801842 A 9801843 A	2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	5-08-2006 1-12-1999 1-12-1999 1-12-1998 1-07-2001 1-09-1998 1-09-1998 1-01-2003 1-05-2000 1-05-2000 1-02-2001 1-0202 1-10-2002 1-10-2002 1-10-2000 1-10-200 1
EP 0492087	Α	01-07-1992	AT DE DE EP	117352 T 4138417 A1 59104330 D1 0492087 A2	15- 11- 02-	-01-1999 -02-1995 -06-1992 -03-1995 -07-1992
DD 296474	Α	05-12-1991	DD	296474 A5		12-1991
US 4623390	Α	18-11-1986	NONE			
ure details about th	B ODDOVI DO Off		*	ent Office, No. 12/82		